

ALL DAY ALLERGY RELIEF- cetirizine hcl capsule
QUALITY CHOICE (Chain Drug Marketing Association)

Cetirizine Hydrochloride Tablets

Active Ingredient (in each capsule)

Cetirizine HCl 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warning

Do not use

if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine.

Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you are

taking tranquilizers or sedatives.

When using this product

- drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding

- if breast-feeding: not recommended
- if pregnant: ask a health professional before use.

Keep out of reach of children

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

Adults and children 6 years and over	One 10 mg capsule once daily; do not take more than one 10 mg capsule in 24 hours. A 5 mg product may be appropriate for less severe symptoms.
Adults 65 years and over	ask a doctor.
Children under 6 years of age	ask a doctor
Consumers with liver or kidney disease	ask a doctor

Other Information

- store at 20° to 25°C (68° to 77° F)
- avoid high humidity and excessive heat above 40°C (104°F)
- protect from light

Inactive Ingredients

Butylated hydroxytolene, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol 400, purified water, sodium hydroxide, Sorbitan, Sorbitol

Questions or comments?

Call 1-248-449-9300 Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the Active Ingredient in ZYRTEC®

All Day Allergy Relief

Indoor & Outdoor Allergies

Cetirizine HCl Capsules, 10 mg

Antihistamine

24 HOUR RELIEF OF:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

Softgels**

(**Liquid-Filled Capsules)

*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Zyrtec®.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARD FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.


Distributed by: C.D.M.A., Inc.©

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Product Label



NDC 63868-432-25


Compare to the Active Ingredient in ZYRTEC


All Day Allergy Relief

Indoor & Outdoor Allergies

Cetirizine HCl Capsules, 10 mg Antihistamine

24 HOUR RELIEF OF:
Sneezing
Runny Nose
Itchy, Watery Eyes
Itchy Throat or Nose





Actual Size

25 Softgels**
(**Liquid-Filled Capsules)

Drug Facts

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Ask a doctor or pharmacist before use if you are taking tranquilizers or sedatives.
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Drug Facts (continued)

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QUALITY CHOICE All Day Allergy Relief

ALL DAY ALLERGY RELIEF

cetirizine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-432
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	

BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)

Product Characteristics

Color	yellow	Score	no score
Shape	OVAL	Size	13mm
Flavor		Imprint Code	10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-432-25	1 in 1 BOX	06/30/2018	02/23/2024
1		25 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207235	06/30/2018	02/23/2024

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 11/2021

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